

Contents lists available at ScienceDirect

Journal of Bone Oncology

journal homepage: www.elsevier.com/locate/jbo

Research Paper

Socioeconomic and demographic factors contributing to outcomes in patients with primary lymphoma of bone

Andrew J. Jacobs^a, Ryan Michels^b, Joanna Stein^c, Adam S. Levin^{b,*}^a Hofstra North Shore-LIJ School of Medicine, 500 Hofstra University, Hempstead, NY 11549, USA^b Department of Orthopaedics, North Shore Long Island Jewish Medical Center, 270-05 76th Avenue, New Hyde Park, NY 11040, USA^c Biostatistics Unit, Feinstein Institute for Medical Research, 350 Community Drive, Manhasset, NY 11030, USA

ARTICLE INFO

Article history:

Received 24 November 2014

Accepted 27 November 2014

Available online 26 December 2014

Keywords:

Lymphoma

Bone

SEER

Socioeconomic

ABSTRACT

Background: Primary lymphoma of bone (PLB) is a rare disease, comprising a malignant lymphoid infiltrate of bone. The goal of this study was to identify socioeconomic, demographic, and anatomic factors as prognostic indicators of survival for this disease using the Surveillance, Epidemiology, and End Results (SEER) database.

Methods: The SEER database was used to identify a study population of 692 patients diagnosed with PLB in the United States from 1989 to 2003. Survival was analyzed using the Kaplan–Meier method, with effects of potential prognostic factors on survival analyzed using the log-rank test. Multivariable analysis was performed by Cox proportional hazards regression.

Results: The overall 5-year survival rate was 49.6%, with a 10-year survival rate of 30.2%. Median overall survival was 4.9 years (95% CI: 3.9, 6.1). In multivariable analysis, age ($p < 0.0001$), marital status ($p = 0.006$), and appendicular vs. axial tumor location ($p = 0.004$) were found to be independent predictors of survival.

Conclusions: This population-based study of PLB identified age, marital status, and tumor location as independent indicators of prognosis. This finding supports the clinical suspicion that an appendicular tumor location confers a better prognosis than an axial tumor location.

© 2015 The Authors. Published by Elsevier GmbH. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

1. Introduction

The rare disease now known as primary lymphoma of bone (PLB) was initially described by Oberlin [1]. Parker and Jackson later described PLB as a distinct entity [2], which is understood to be a malignant lymphoid infiltrate of bone [3]. While this may include cortical or soft tissue invasion, the diagnosis generally excludes lymph node or distant visceral involvement to be considered a primary lymphoma of bone [4]. This definition has been contended throughout the literature, with some authors permitting lymph node involvement [5–7], though most authors recognize the disease as lymphoma localized only to bone upon diagnosis.

Primary lymphoma of bone affects 1.7/1,000,000 patients in the US [8]. Previous studies have suggested PLB comprise 5% of all extranodal lymphomas [9], and 3% of all bone malignancies [10].

* Corresponding author. Present address: Department of Orthopaedic Surgery, Johns Hopkins University, 601 N. Caroline Street, Baltimore, MD 21287, USA. Tel.: +1 410 502 2698; fax: +1 410 614 1451.

E-mail addresses: ajacob24@pride.hofstra.edu (A.J. Jacobs), rmichels@nshs.edu (R. Michels), jstein4@nshs.edu (J. Stein), alevin25@jhmi.edu (A.S. Levin).

<http://dx.doi.org/10.1016/j.jbo.2014.11.002>

2212-1374/© 2015 The Authors. Published by Elsevier GmbH. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Due to the low incidence of PLB, most prior studies have been from a single-institution, and have had small sample sizes [10–15]. The Surveillance, Epidemiology, and End Results (SEER) database is a publically available database created for the purpose of collecting demographic, clinical, and outcome data for cancer patients in the United States. There remains scant literature published using this database to evaluate larger patient populations [8].

The purpose of this population-based study was to identify prognostic factors for survival in patients with PLB, in order to determine whether disparities in survival exist between demographic cohorts. In doing so, the current study aimed to determine the prognostic significance of different tumor-specific and anatomic considerations that may influence overall survival from this disease. The authors chose to limit this study to adult patients, because PLB in children is often considered and treated as a systemic disease [16].

2. Material and methods

The study population was selected from the National Cancer Institute's SEER database. The SEER database collects data from 18 geographic registries, covering approximately 28% of the U.S.

population [17]. The SEER*Stat software (Version 8.0.4; NCI; Bethesda, MD) was used to identify 997 adult patients diagnosed with primary lymphoma of bone during a 15-year period from 1989 to 2003. Histology was selected by using ICD-O-3 codes 9590/3, 9591/3, 9670/3, 19671/3, 9675/3, 9680/3, and 9684/3. Primary site was selected as C40.0, C40.1, C40.2, C40.3, C40.8, C40.9, C41.2, C41.3, C41.4, and C41.9. Exclusion criteria included lesions of the skull and face, T-cell lymphoma, and cases without follow up data, yielding a final study population of 692 patients.

Tumor location was dichotomized as either appendicular or axial. The scapula was considered to be part of the appendicular skeleton, while the pelvic bones were considered to be part of the axial skeleton. Marital status was categorized as single, married, or other (including separated, divorced or widowed). Rural–urban continuum code was collapsed into a binary variable: Metro county or non-metro county, using guidelines by SEER and the Economic Research Service [18,19]. SEER registry region was aggregated into regions (Northeast, South, Southwest, Midwest, and West). Race was categorized as White, Black or Asian/Other. Age was considered as a categorical variable (< 30 years, 30–59 years, ≥ 60 years).

Statistical analysis was performed in SAS version 9.3 (SAS Institute, Cary, NC). The effects of categorical variables on survival were assessed by computing Kaplan–Meier product limit curves and compared using the log-rank test. The effects of continuous variables were analyzed using Cox proportional hazards regression. The Bonferroni method was applied when performing multiple comparisons. Factors that appeared to be significantly associated with survival in the univariate analysis were considered for inclusion in the final multivariable Cox proportional hazards regression model. A result was considered statistically significant with a *p*-value < 0.05. Efron's method was used to adjust for tied failure times.

Incidence rates were age-adjusted to the 2000 US standard population, with confidence intervals calculated using the Tiwari modification. Annual percentage change was calculated using the weighted least squares method.

3. Results

The final analysis included 692 patients, whose demographic and clinical characteristics are presented in Table 1. The majority of patients were white (89.0%), non-Hispanic (91.3%), and lived in metropolitan counties (87.4%). The majority of patients were over the age of 60 years (55.6%), and diffuse large B-cell lymphoma was the most common histologic classification (71.2%). The western region of the United States contributed the largest proportion of patients to the study population (56.5%).

The estimated overall survival of patients for all patients in this study was 49.6% at 5-years, and 30.2% at 10-years (Fig. 1). The incidence of PLB during the 15-year study period ranged from 0.1/100,000 to 0.3/100,000 (Fig. 2). The annual percent change for this time period was non-significant, suggesting a stable incidence over the study period.

In univariate analysis, significant factors for overall survival included age ($p < 0.0001$), marital status ($p < 0.0001$), anatomic location of tumor ($p < 0.0001$), geographic region ($p = 0.02$), and tumor grade ($p = 0.01$). After Bonferroni adjustment, tumor grade was no longer a significant prognostic indicator for overall survival. Furthermore, after Bonferroni adjustment for multiple comparisons, tumor grade and geographic region were not statistically significantly associated with overall survival. Univariate analysis results for categorical variables are presented in Table 2. Kaplan–Meier product limit curves are provided for age (Fig. 3), marital status (Fig. 4), and tumor location (Fig. 5).

Table 1

Descriptive demographic and clinical statistics of the study population.

Characteristic	Frequency	% Of total
Total number of patients	692	100.0
Age		
< 30	67	9.7
30 to 59	240	34.7
60 or greater	385	55.6
Sex		
Male	370	53.5
Female	322	46.5
Race		
White	616	89.0
Black	45	6.5
Asian, other and unknown	31	4.5
Ethnicity		
Non-Spanish–Hispanic–Latino	632	91.3
Spanish–Hispanic–Latino	60	8.7
Marital status		
Single	122	18.2
Married	383	57.2
Divorced, separated, widowed	165	24.6
Geographic region		
Southeast	54	7.8
South	42	6.1
Midwest	111	16.0
Northeast	94	13.6
West	391	56.5
County		
Non-metro county	87	12.6
Metro county	605	87.4
Tumor site		
Axial	389	56.2
Appendicular	303	43.8
Histology		
Malignant lymphoma, NOS ^a	46	6.7
Malignant lymphoma, non-Hodgkin, NOS ^a	75	10.8
Malignant lymphoma, small B lymphocytes, NOS ^a	21	3.0
Lymphoplasmacytic lymphoma (NHL ^b)	11	1.6
Malignant lymphoma, mixed small and large cell, diffuse	22	3.2
Diffuse large B-cell (NHL ^b) lymphoma	493	71.2
Malignant lymphoma, large B, diffuse, immunoblastic	24	3.5
Grade		
B-cell; pre-B; B-precursor	668	96.5
Well differentiated; grade I	6	0.9
Moderately differentiated; grade II	9	1.3
Poorly differentiated; grade III	5	0.7
Undifferentiated; anaplastic; grade IV	4	0.6
Surgery		
Yes	177	25.8
No	510	74.2
Radiation		
Yes	475	70.6
No	198	29.4

^a NOS, not otherwise specified.

^b NHL, non-Hodgkin lymphoma.

The final multivariable model demonstrated that age ($p < 0.0001$), marital status ($p = 0.02$), and appendicular or axial tumor location ($p = 0.004$) remained significant independent prognostic variables for overall survival (Table 3). A survival advantage was demonstrated for younger patients. The mortality rate for PLB patients in the 30–59 age group is estimated to be 4.4 times that for those patients in the < 30 age group, after adjusting for marital status and tumor location (CI: 1.7–11.2; $p = 0.002$). Furthermore, patients aged 60 or older are estimated to have a mortality rate 12.8 times that for < 30 year-old

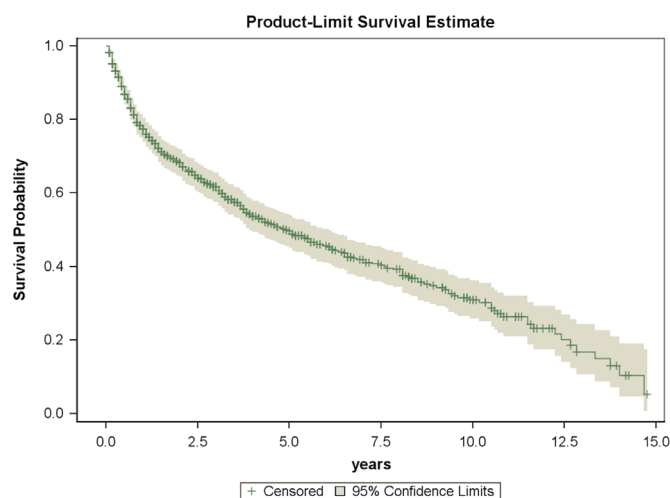


Fig. 1. Kaplan-Meier plot of overall survival in study population.

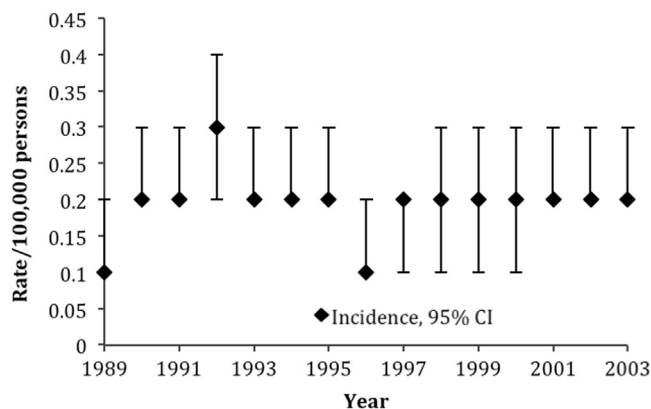


Fig. 2. Incidence of primary lymphoma of bone, age-adjusted to the 2000 US standard population.

patients, after adjusting for tumor location and marital status (CI: 5.1–32.3; $p < 0.0001$), though this finding is potentially influenced by medical comorbidities.

The multivariable Cox regression model also showed that patients who were divorced, separated, or widowed had a significantly worse outcome than patients who were married or single; the mortality rate for Divorced, Separated or Widowed PLB patients was estimated to be 1.4 times that for married patients, after adjusting for tumor location and age (CI: 1.1–1.8; $p = 0.006$). The hazard ratio for comparing single with married patients was non-significant, however ($p = 0.69$).

Anatomical location of the primary tumor was also a significant factor in overall survival. Patients with an appendicular lesion had significantly better prognosis than patients with a primary axial lesion (Fig. 6). The mortality rate for PLB patients with axial-located disease is estimated to be 1.4 times that for those patients with appendicular-located disease, after adjusting for marital status and age (CI: 1.2–1.8; $p = 0.004$).

4. Discussion

The current study presents a population-based analysis of the effects of demographic factors on overall survival from primary lymphoma of bone. In order to account for the various clinical indicators, this study utilized the large patient population to better analyze the relative prognostic significance of patient- and tumor-related predictors of overall survival from the disease. To our knowledge, this is one of the largest studies of patients with primary lymphoma of bone published in the English language medical literature. Among the strengths of a large, population-based study is the inclusion of a larger collection of patients than would be feasible for a single institution study of such a rare disease, as well as the ability to sample a cross-section of the United States' population.

Single institution studies have been published on individual chemotherapy regimens, as well as the individual prognostic

Table 2
Univariate analysis of effects on overall survival.

Characteristic	1-Year survival	5-Year survival	10-Year survival	p-Value ^a
Overall	77.4	49.6	30.2	NA ^b
Age				< .0001*
< 30	98.5	NA	NA	
30–59	85.5	71.8	52.7	
> 60	68.8	30.6	13.9	
Sex				.24
Male	78.0	49.5	35.6	
Female	76.7	49.8	24.5	
Race				.09
White	77.2	48.2	29.4	
Black	82.8	62.2	NA	
Other	72.8	63.4	50.7	
Ethnicity				.40
Non-Hispanic	78.2	50.0	30.7	
Hispanic	65.3	45.9	17.8	
Marital status				< .0001*
Married	79.5	50.5	33.9	
Single	86.0	69.0	50.0	
Divorced, separated, widowed	66.6	34.3	11.6	
Geographic region				.02*
Northeast	77.4	50.3	NA	
Southeast	73.1	8.1	0	
Midwest	73.8	46.3	26.1	
West	77.6	54.0	34.9	
South	78.8	41.7	34.2	

Table 2 (continued)

Characteristic	1-Year survival	5-Year survival	10-Year survival	p-Value ^a
Tumor location				<.0001*
Appendicular	85.6	58.4	33.7	
Axial	71.1	43.1	28.4	
Histology				1.0
Malignant lymphoma, NOS ^c	78.0	46.7	32.4	
Malignant lymphoma, non-Hodgkin, NOS ^c	73.2	39.2	NA	
Malignant lymphoma, small B lymphocytes, NOS ^c	69.7	25.7	NA	
Lymphoplasmacytic lymphoma (NHL ^d)	81.8	61.4	NA	
Malignant lymphoma, mixed small and large cell, diffuse	80.6	65.1	18.3	
Diffuse large B-cell (NHL ^d) lymphoma	77.9	50.4	31.5	
Malignant lymphoma, large B, diffuse, immunoblastic	66.7	51.9	28.8	
Grade				.24
B-cell; pre-B; B-precursor	77.6	50.4	31.1	
Grade I	66.7	NA	NA	
Grade II	77.8	38.9	NA	
Grade III	60.0	NA	NA	
Grade IV	NA	NA	NA	
Surgery				0.19
Yes	78.6	42.1	27.4	
No	77.1	52.8	32.3	
Radiation				0.79
No	73.2	50.0	29.5	
Yes	80.3	49.2	29.9	

^a p Values are results of the log-rank test, with Bonferroni adjustments made for multiple comparisons.

^b NA, not available.

^c NOS, not otherwise specified.

^d NHL, non-Hodgkin lymphoma

* Indicates result is statistically significant on a 0.05 level of significance.

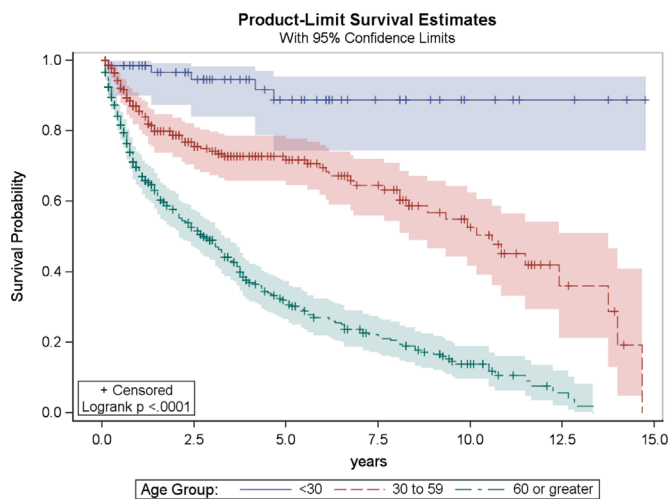


Fig. 3. Kaplan-Meier plot of overall survival related to age at diagnosis.

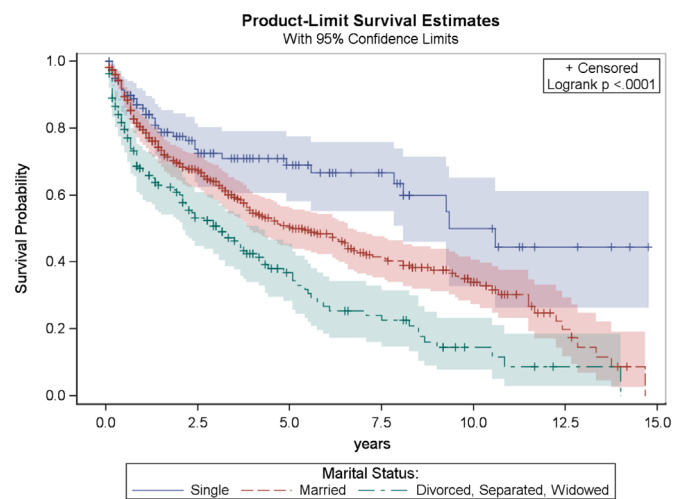


Fig. 4. Kaplan-Meier plot of overall survival related to marital status.

potentials of several laboratory values (i.e. Lactate Dehydrogenase) [10,15]. However, the SEER database does not include data on chemotherapy [17], thus the relative effect of chemotherapy treatment regimens could not be evaluated in our analysis. We addressed the concern over evolving chemotherapy protocols by limiting the study to a 15-year period. While the authors recognize that significant improvements in imaging, staging, and clinical care have been made over the study period, there was no significant change in the incidence or overall survival over the course of this study, thereby supporting a stable clinical period for analysis. Limiting the years of the study also helped control for the changes in classification and categorization of the disease, while allowing for long-term follow-up for overall survival.

We found that appendicular tumor location is an independent predictor of survival in multivariable analysis when compared with axial location. While this has been suggested by previous authors, other relatively large studies investigating PLB have not yielded significant prognostic importance for overall survival, when considering confounding variables [8,15]. Age at time of diagnosis was also an independent predictor of survival, which has been consistently demonstrated in prior analyses [8,13,14]. Age over 60 years has previously been shown to influence both overall and disease-specific survival, so these findings are not entirely related to mortality from unrelated conditions. However, the ability to endure chemotherapy due to age and comorbidities may play a significant role in considering treatment options.

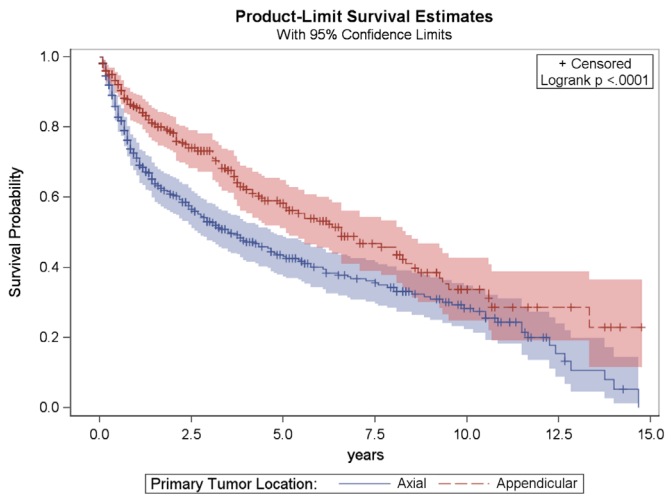


Fig. 5. Kaplan-Meier plot of overall survival related to primary tumor location.

Table 3
Multivariable analysis results.

Variable	Hazard ratio	95% CI ^a	p-Value
Tumor location			
Appendicular		Reference group	
Axial	1.41	1.12–1.77	0.0035*
Marital status			
Married		Reference group	
D, S, W ^b	1.41	1.11–1.79	0.0055*
Single	1.08	0.75–1.56	0.6889
Age			
< 30		Reference group	
30 to 59	4.39	1.72–11.19	0.0019*
60 or greater	12.81	5.08–32.29	< .0001*

^a CI, confidence interval.
^b D, divorced; S, separated; W, widowed.
* Indicates statistically significant on a $p < 0.05$ level.

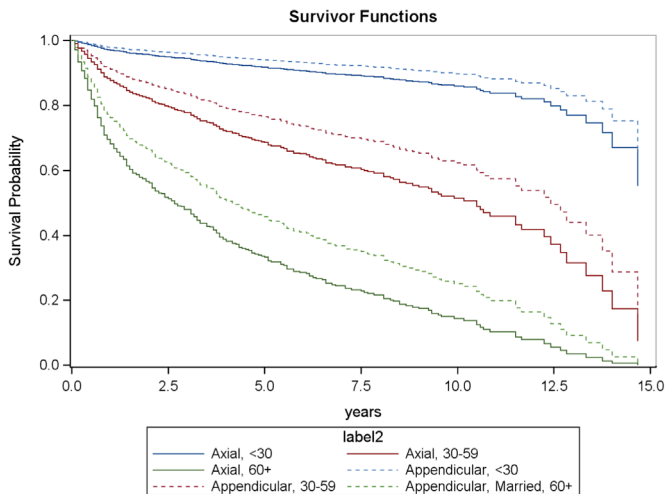


Fig. 6. Multivariate adjusted survival curve comparing tumor location from 1989 to 2003 among married patients, stratified by age.

In the current study, multivariable analysis demonstrated that divorced, separated, or widowed patients have a significantly

poorer prognosis than married patients, independent of age. There was insufficient data available from the SEER database to elucidate the cause of this finding. However, using the SEER database, Aizer et al., also demonstrated that marital status was an independent predictor of survival following oncologic diagnoses [20]. They reported that the survival benefit associated with marriage was larger than the published survival benefit of chemotherapy for prostate, breast, colorectal, esophageal and head/neck cancers. Marital status was associated with lower stage at presentation in that study [20], suggesting that these findings may be due to lead time bias of earlier detection in married patients, rather than a true reflection of improved mortality.

5. Conclusions

In conclusion, there is evidence to suggest that socioeconomic factors, as well as clinical factors, contribute to the overall survival in patients with primary lymphoma of bone. Younger age, appendicular tumor location, and being married appear to be good prognostic factors for survival among patients with PLB.

Conflict of interest statement

The authors declare that there are no conflicts of interest.

References

[1] Oberlin C. Les reticulosarcomes at les reticuloenotheliosarcomes de la moelle osseuse sarcomes d'Ewing. Bull Assoc Fr Etud Cancer 1928;17:259–96.

[2] Parker J, Jackson H. Primary reticulum cell sarcoma of bone. Surg Gynecol Obstet 1939;68:45–53.

[3] Pettit CK, et al. Primary lymphoma of bone. A B-cell neoplasm with a high frequency of multilobated cells. Am J Surg Pathol 1990;14(4):329–34.

[4] Boston Jr HC, et al. Malignant lymphoma (so-called reticulum cell sarcoma) of bone. Cancer 1974;34(4):1131–7.

[5] Ferreri AJ, et al. Therapeutic management with adriamycin-containing chemotherapy and radiotherapy of monostotic and polyostotic primary non-Hodgkin's lymphoma of bone in adults. Cancer Invest 1998;16(8):554–61.

[6] Shoji H, Miller TR. Primary reticulum cell sarcoma of bone. Significance of clinical features upon the prognosis. Cancer 1971;28(5):1234–44.

[7] Glotzbecker MP, et al. Primary non-Hodgkin's lymphoma of bone in children. J Bone Joint Surg Am 2006;88(3):583–94.

[8] Jawad MU, et al. Primary lymphoma of bone in adult patients. Cancer 2010;116(4):871–9.

[9] Freeman C, Berg JW, Cutler SJ. Occurrence and prognosis of extranodal lymphomas. Cancer 1972;29(1):252–60.

[10] Beal K, Allen L, Yahalom J. Primary bone lymphoma: treatment results and prognostic factors with long-term follow-up of 82 patients. Cancer 2006;106(12):2652–6.

[11] Durr HR, et al. Malignant lymphoma of bone. Arch Orthop Trauma Surg 2002;122(1):10–6.

[12] Ford DR, et al. Primary bone lymphoma—treatment and outcome. Clin Oncol (R Coll Radiol) 2007;19(1):50–5.

[13] Horsman JM, et al. Primary bone lymphoma: a retrospective analysis. Int J Oncol 2006;28(6):1571–5.

[14] Lewis VO, et al. Oncologic outcomes of primary lymphoma of bone in adults. Clin Orthop Relat Res 2003(415):90–7.

[15] Demircay E, et al. Malignant lymphoma of bone: a review of 119 patients. Clin Orthop Relat Res 2013;471(8):2684–90.

[16] Doll C, et al. Primary B-cell lymphoma of bone in children. Eur J Pediatr 2001;160(4):239–42.

[17] Surveillance Epidemiology and End Results. [accessed: July 30, 2013]; available from (<http://seer.cancer.gov/>); 2012.

[18] Rural Classifications. [accessed: July 30, 2013]; Available from: (<http://www.ers.usda.gov/topics/rural-economy-population/rural-classifications.aspx>); 2013.

[19] Rural-Urban Continuum Codes. [accessed: July 30, 2013]; Available from: (<http://seer.cancer.gov/seerstat/variables/countyattribs/ruralurban.html>); 2012.

[20] Aizer AA, et al. Marital status and survival in patients with cancer. J Clin Oncol 2013;31(31):3869–76.